CLAIMS

- 1. Method for obtaining, preparing or producing human suppressor T lymphocytes (and/or the precursors thereof), comprising a step of selection, separation or isolation in vitro or ex vivo of human T lymphocytes expressing the THY-1 molecule.
- 2. Method according to claim 1, comprising:

5

10

25

- (a) obtaining a cell population of human origin comprising T lymphocytes, and
- (b) recovering T lymphocytes expressing the THY-1 antigen.
- 3. Method according to claim 1, characterized in that step (b) is preceded or followed by a step of amplification of T lymphocytes.
- 4. Method according to any one of the previous claims, characterized in that the T lymphocytes expressing the THY-1 antigen are selected, separated, isolated or recovered by means of a ligand specific of THY-1.
- 5. Method according to claim 4, characterized in that the specific ligand is an antibody specific of THY-1 or a fragment or derivative of said antibody having substantially the same antigenic specificity.
 - 6. Method according to claim 5, characterized in that the specific ligand is a monoclonal or polyclonal antibody specific of THY-1.
 - 7. Method according to claim 5, characterized in that the specific ligand is a polyfunctional, monocatenary or multimeric antibody, specific of THY-1.
 - 8. Method according to claim 4, characterized in that the specific ligand is an aptamer.

- 9. Method according to any one of claims 4 to 8, characterized in that the ligand is immobilized on a support or placed in solution.
- 10. Method according to claim 9, characterized in that the support is a column or a bead, preferably a magnetic bead.
- 11. Method according to any one of claims 4 to 10, characterized in that the ligand is labelled.
- 12. Method according to claim 11, characterized in that the labelling is carried out by means of a fluorescent, radioactive, luminescent, phosphorescent, chemical or enzymatic detection label.
 - 13. Method according to any one of the previous claims, characterized in that the step of recovery, selection or isolation is carried out by flow cytometry, affinity chromatography, FACS, MACS or D/MACS.
 - 14. Method according to any one of the previous claims, characterized in that the cell population comes from a tissue selected in the group consisting of bone marrow, spleen, liver, thymus, blood which has or has not been previously enriched in T lymphocytes, umbilical cord blood, fetal, infant or adult peripheral blood, a tumor, a site of inflammation, a transplanted organ or a cell culture established with one or another of said tissues.
- 25 15. Method for identifying and/or quantifying human suppressor T lymphocytes in a cell population, comprising exposing said cell population to a ligand specific of THY-1 and determining and/or quantifying the formation of a complex between the ligand and the cells, the formation of said complexes indicating the presence and/or the quantity of suppressor T lymphocytes in the cell population.

30

5

15

- 16. Method for producing a pharmaceutical composition, comprising:
 - (a) obtaining a biological sample comprising human T lymphocytes,

- (b) selecting T lymphocytes expressing the THY-1 antigen in said biological sample, and
- (c) conditioning said T lymphocytes expressing the THY-1 antigen in a pharmaceutically acceptable adjuvant or medium.

5

- 17. Method for producing a pharmaceutical composition, comprising:
 - (a) obtaining a biological sample comprising human T lymphocytes,
 - (b) depleting T lymphocytes expressing the THY-1 antigen from said biological sample, and

10

- (c) conditioning said T lymphocytes not expressing the THY-1 antigen in a pharmaceutically acceptable adjuvant or medium.
- 18. Use of a ligand specific of the THY-1 antigen in order to select, identify, sort or prepare, in vitro or ex vivo, human suppressor T lymphocytes.

15

- 19. Use of a ligand specific of the THY-1 antigen in order to prepare a diagnostic composition intended for the selection, identification or quantification in vivo of human suppressor T lymphocytes.
- 20. Use of a ligand specific of the THY-1 antigen in order to prepare a therapeutic composition intended for the modification, stimulation or elimination in vivo of human suppressor T lymphocytes.
 - 21. Isolated human T lymphocyte, characterized in that it exhibits suppressor activity and in that it expresses the CD8 or CD4 markers and THY-1.
 - 22. Cell composition comprising at least 50, 60, 70, 80, 85, 90 or 95% of human CD8+/THY-1+ or CD4+/THY-1+ T cells.
- 30 23. T lymphocyte according to claim 21, characterized in that it is genetically modified by means of a viral vector.

- 24. Pharmaceutical composition characterized in that it comprises at least one suppressor T lymphocyte according to any one of claims 21 to 23 and a pharmaceutically acceptable adjuvant or medium.
- 5 25. Use of a T lymphocyte or a population of T lymphocytes according to any one of claims 21 to 23 for preparing a composition intended to implement a therapeutic method.

10

15

- 26. Use according to claim 25, characterized in that the composition is intended to be used as a vaccine.
- 27. Use according to claim 25 for preparing a composition intended for the treatment of a tumor, an autoimmune disease, an allergy, graft-versus-host disease, an inflammatory disease, type 1 diabetes, a viral or bacterial infection, for immune reconstitution or for induction of tolerance in the event of stem cell, tissue or organ transplantation in a mammal.
- 28. Kit for the isolation or characterization of human suppressor T lymphocytes comprising a ligand specific of THY-1, placed in solution or on a support, and, optionally, reagents for detection of the ligand.